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IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re application of : **Confirmation No. 7206**
Yasuhiro TADA et al. : Attorney Docket No. 2006_0371A
Serial No. 10/574,111 : Group Art Unit 1797
Filed March 31, 2006 : Examiner Ana M. Fortuna

VINYLIDENE FLUORIDE BASED RESIN
POROUS HOLLOW YARN AND
METHOD FOR PRODUCTION THEREOF : **Mail Stop: Amendment**

SUBMISSION OF RULE 132 DECLARATION

Commissioner for Patents
P.O. Box 1450
Alexandria, VA 22313-1450

THE COMMISSIONER IS AUTHORIZED
TO CHARGE ANY DEFICIENCY IN THE
FEE FOR THIS PAPER TO DEPOSIT
ACCOUNT NO. 23-0975.

Sir:

On June 11, 2008 Applicants filed an Amendment in response to the Office Action of December 11, 2007.

In further support of the patentability of the presently claimed invention, Applicants are submitting herewith a Rule 132 Declaration which demonstrates the importance of stretching the layer of a vinylidene fluoride resin, and the use of a specific blend of types of polyvinylidene fluoride resins having an ultra-high molecular weight and a medium-to-high molecular weight in accordance with the present invention.

Respectfully submitted,

Yasuhiro TADA et al.

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In re Application of :
Yasuhiro TADA et al.
Application S.N.: 10/574,111
Filed: March 31, 2006

Group Art Unit: 1797
Examiner: Ana M. Fortuna

For: VINYLIDENE FLUORIDE BASED
RESIN POROUS HOLLOW YARN
AND METHOD FOR PRODUCTION
THEREOF

Assistant Commissioner for Patents
Washington, D.C. 20231

Sir:

DECLARATION UNDER 37 CFR 1.132

I, the undersigned, Yasuhiro TADA, hereby declare as follows:

1. I am a citizen of Japan and a resident of 18 -13 Kamitamari, Omitama-shi, Ibaraki-ken 311-3436, Japan.
2. In March 1991, I received my Master of Engineering degree in polymer material science from the Faculty of Science and Engineering of Yamagata University.
3. Since April 1991, I have been employed at Kureha Corporation (formerly, Kureha Kagaku Kogyo Kabushiki Kaisha) and have conducted research and development in the filed of, among others, stretched polychlorotrifluoroethylene film and polyvinylidene fluoride hollow fiber membrane, in Polymer Processing & Products Research Laboratories of Kureha Corporation.
I am an inventor of U.S. Patent No. 5,833,070 regarding stretched polychlorotrifluoroethylene film and U.S. Patent No. 7,351,338 regarding polyvinylidene fluoride hollow fiber membrane.
4. I am one of the applicants of the application S.N.: 10/574,111

(hereinafter referred to as the instant application) and accordingly I am familiar with the specification and claims of the instant application.

5. Moreover, I have read carefully and I am familiar with the Official Action dated December 11, 2007, which action rejected Claims 1 – 8 of the instant application. I have read carefully and I am familiar with several references, inclusive of Takamura et al (US 6, 299,773), Nohmi et al (US 4,399,035) and Morikawa et al (US 7,258,914) which are hereinafter referred to as Takamura ('773), Nohmi ('035) and Morikawa('914), respectively. The substance of the Examiner's rejection from the prior art aspect is believed to be based on the conclusion that Claims 1 – 7 are anticipated by or, in the alternative obvious over Takamura ('773); claims 1 – 3 and 6 are obvious over Nohmi ('035) or Morikawa('914).

6. In view of the Examiner's rejection, the applicants have amended claim 1 of the instant application based on the features of claims 5 and 8 as filed of the instant application to recite: that the claimed porous hollow fiber comprises a stretched single layer of a vinylidene fluoride resin; and that the vinylidene fluoride resin comprises 2-49 wt.% of a first vinylidene fluoride resin having a weight-average molecular weight (Mw_1) of 4×10^5 - 12×10^5 and 51-98 wt.% of a second vinylidene fluoride resin having a weight-average molecular weight (Mw_2) of 1.5×10^5 - 6×10^5 provided that the weight-average molecular weight (Mw_1) of the first vinylidene fluoride resin and the weight-average molecular weight (Mw_2) of the second vinylidene fluoride resin satisfy a ratio Mw_1/Mw_2 of at least 1.2.

7. For demonstrating the importance of the stretching and the use of a specific blend of two types of polyvinylidene fluoride (PVDF) having an ultra-high molecular weight and a medium-to-high molecular weight not disclosed by any of Takamura ('773), Nohmi ('035) and Morikawa('914), I made, under my direction and control some experimental tests, the procedure and the results of which are reported hereinbelow.

<EXPERIMENTS>

(Comparative Example A)

The production of a hollow fiber membrane was tried by using a single species of polyvinylidene fluoride (PVDF) having a weight-average molecular

weight of 4.90×10^5 instead of the PVDF used in Example 7 otherwise in a similar manner as in Example 7 of the instant application. In the comparative test, however, a nozzle having a nozzle having an annular slit of 6 mm in outer diameter and 4 mm in inner diameter was used instead of the nozzle having an annular slit of 5 mm in outer diameter and 3.5 mm in inner diameter because of unavailability of the latter nozzle due to optimization of the nozzle after the instant application.

More specifically, a first intermediate form (i.e., a hollow fiber before extraction of the plasticizer) was prepared in the same manner as in Example 7 of the instant application except that the nozzle having an annular slit of 6 mm in outer diameter and 4 mm in inner diameter was used, and correspondingly, air was injected into a hollow part of the extruded fiber at a rate of 4.7 ml/min. and the extruded and cooled fiber was taken up at a rate of 3m/min., wherein the latter two modifications were adopted so as to provide a first intermediate form having an outer diameter of 1.943 mm and an inner diameter of 1.374 comparable to those if the first intermediate form of Example 7.

The first intermediate form was then subjected to extraction with dichloromethane, drying to remove the dichloromethane and heat treatment in the same manner as in Example 7 to prepare a second intermediate form (i.e., an unstretched hollow fiber membrane).

The second intermediate form was longitudinally stretched at a ratio of 1.35 times (substantially lower than 1.7 times in Example 7) at an environmental temperature of 25 °C, whereas the fiber was severed. Accordingly, the longitudinal stretch ratio was lowered to 1.30 times to obtain a stretched hollow fiber membrane, which was then subjected to two times of immersion in dichloromethane and heat fixation in the same manner as in Example 7 to obtain a stretched porous hollow fiber membrane.

The properties of the unstretched and stretched hollow fiber membranes were evaluated in the same manner as described in the instant application. The outline of the production conditions and the measured properties were shown in the attached Table A

In Table A, the outline of the production conditions and the measured properties of the stretched hollow fiber membrane of Example 7 shown in Table 1 of the instant application and the stocked data of the corresponding unstretched hollow fiber membrane, are also included for convenience of comparison.

<EVALUATION>

In view of the results shown in Table A, the following evaluation is believed to be readily derivable.

- (1) From a comparison between Example 7 and Example 7(Unstretched), a remarkable increase in basic water permeability as an essential performance of a hollow fiber membrane for water treatment is attained can be attained as a result of stretching.
- (2) From a comparison between Example 7 and Comparative Example A, the use of a specific blend of two types of polyvinylidene fluoride (PVDF) having an ultra-high molecular weight and a medium-to-high molecular weight remarkably facilitate the stretching of a yet-unstretched hollow fiber membranes.

I further declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true, and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issuing thereon.

Dated: July 4, 2008

Yasuhiro Tada
Yasuhiro TADA

Attachment: Table A

Table A :

EXAMPLE		7	7	Comparative A (Unstretched)	Comparative A (x1.35 stretch)	Comparative A (x1.30 stretch)	Comparative A (Unstretched)
Starting material composition	Mixture A	1st. PVDF's Mw (Mw1) ($\times 10^5$)	9.36	9.36	-	-	-
		2nd. PVDF's Mw (Mw2) ($\times 10^5$)	4.12	4.12	4.90	4.90	4.90
		Mw1/Mw2	2.27	2.27	-	-	-
		1st. PVDF/2nd. PVDF mixing ratio (wt. %)	5/95	5/95	-	-	-
		Mixture's Mw ($\times 10^5$)	4.38	4.38	-	-	-
		Mw/Mn	-	-	-	-	-
	Mixture B	Polyester plasticizer	PN-150	PN-150	PN-150	PN-150	PN-150
		Solvent	NMP	NMP	NMP	NMP	NMP
		Plasticizer/solvent mixing ratio (wt. %)	82.5/17.5	82.5/17.5	82.5/17.5	82.5/17.5	82.5/17.5
Spinning and stretching conditions	Mixture A/Mixture B Supply ratio (wt. %)		35.7/64.3	35.7/64.3	35.7/64.3	35.7/64.3	35.7/64.3
	Nozzle O. D. (mm)		5	5	6	6	6
	Nozzle I. D. (mm)		3.5	3.5	4	4	4
	Air supply rate (ml/min)		6.2	6.2	4.7	4.7	4.7
	Air gap (mm)		170	170	170	170	170
	Water bath temp. (°C)		60	60	60	60	60
	Take-up speed (m/min)		5	5	3	3	3
	Do before extraction (mm)		1.949	1.949	1.943	1.943	1.943
	Di before extraction (mm)		1.378	1.378	1.374	1.374	1.374
	Stretch ratio (times)		1.7	1.00	1.35	1.30	1.00
	Outer diameter Do (mm)		1.57	1.697	*3	1.453	1.580
	Inner diameter Di (mm)		1.072	1.173		1.052	1.147
	Thickness (mm)		0.249	0.262		0.200	0.217
Physical properties	Porosity (%)		75.9	63.3		57.9	54.1
	Ave. pore size P (μ m)		0.131	0.070		0.095	0.055
	Max. pore size (μ m)		0.277	0.153		0.190	0.150
	C value (/day, 100kPa at 25°C)		-8.7	-0.001		-0.01	-0.002
	Water permeability Fo value ($\text{m}^3/\text{m}^2 \cdot \text{day}$, 100kPa at 25°C)		72.2	11.8		50.1	4.8
	Fo/P		551.1	169.5		529.1	87.3
	Fo/Di ⁴		54.7	6.2		41.0	2.8
	Tensile strength (MPa)		10.9	10.2		10.1	5.8
	Elongation at break (%)		18.2	141		9.9	51.2

*3: Hollow fiber was severed during the stretching.